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In response to the Advisory Action of February 18, 2004 in the above-identified application, please amend the application as follows:

IN THE SPECIFICATION

At page 1, please replace the original structural formula with the following:

Page 1, lines 18-19, please insert the following:

X1, X2, X3, X4, same or different, are a group chosen among: -CONR-, -NRCO--CH2-NR-, -NR-CH2- where R is H, C1-3 alkyl, or benzyl;

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Page 1, lines 22-27, please insert the following:

-(CH₂)_r Ar₁ where r is 0, 1 or 2 and Ar is an aromatic group chosen among benzene. naphthalene, thiopene, benzothiophene, pyridine, quinoline, indole, furan, benzofuran, thiazole, benzothiazole, imidazole, benzoimidazole, possibly substituted with up to 2 substituents chosen among C1-3 alkyl, C1-3 haloalkyl, C1-3 alkyloxy and C2-4 amino-alkyloxy, halogens, OH, NH2, CN, NR₆R₇, where R₆ and R₇, are the same or different, and are H or C₁₋₃ alkyl,

At page 2, lines 2-12, please replace with the following:

15 --Ro is a methanesulfonyl, tosyl, tetrahydropyranyl, tetrahydrothiopyranyl possible mono or disubstituted by oxygen on the S atom, piperidyl possibly optionally substituted on the N atom by a C₁₋₃ alkyl, C₁₋₃ acyl, aminosulfonyl or methanesulfonyl; or a group (CH₂)_gR₁₀ where g g is 1,2, or 3 and R₁₀ is chosen among morpholine, furan, or CN; or R₈ and R₉ together with the N atom to which they are linked form a piperazine possibly optionally substituted at the other N atom one of its nitrogen 20 atoms by C₁₋₃ alkyl, C₁₋₃ acyl or methanesulfonyl;--

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At page 2, lines 13-21, please insert the following:

-N(R₁₁)CO(CH₂)_h-R₁₂ where R₁₁ is H $\sqrt{\underline{or}}$ C₁₋₃ alkyl; h is 0, 1, 2 \underline{or} 3; and R₁₂ is chosen among: morpholine, pyrrolidine possibly substituted with an hydroxy or an hydroxymethyl, piperidine possibly substituted with a group hydroxy, carboxyamido or aminosulfonyl, piperazine possibly substituted on the N-atom by C₁₋₃ alkyl, triazole, tetrazole, 5-mercapto-tetrazole, furan, thiophene, thiomorpholine possibly mono or di-oxygenated on the S-atom, and amino-cyclohexane possibly substituted by an hydroxy group f; or

-COR₁₃ wherein R₁₃ is morpholine or piperazine possibly substituted with a C₂₋₆

alkyl containing one or more ether or hydroxy groups/; and Rs is H. 30

At page 4, line 3, please insert:

--R₁ and R₂ same or different, are: --

At page 4, lines 14-27, please replace with the following:

- R9₂ is chosen among: methanesulfonyl, tosyl, tetrahydropyranyl, tetrahydrothiopyranyl possibly mono or di-substituted by oxygen on the S atom, piperidyl possibly substituted on the N-atom by a C1-3₁₋₃ alkyl, C1-3₁₋₃ acyl, aminosulfonyl or methanesulfonyl; or a group (CH2₂)g₂-R₁₀ where g is 1, 2 or 3 and R₁₀ is chosen among morpholine, furan or CN;
- or R₈ and R₉ together with the N atom to which they are linked form a piperazine possibly substituted on the other N atom with a C1-3₁₋₃ alkyl, C1-3₁₋₃ acyl or methanesulfonyl;
 - $-N(R_{11})CO(CH_2)_h-R_{12}$ where R_{11} is H or C_{1-3} alkyl; h is 0, 1, 2, or 3;
 - and R₁₂ is chosen among: morpholine, pyrrolidine possibly optionally substituted with an hydroxy or hydroxymethyl, piperidine possibly optionally substituted with a group 4-hydroxy, or 4-carboxyamido group or aminosulfonyl, piperazine possibly optionally substituted on the other N-
- atom by C₁₋₃ alkyl, triazole, tetrazole, 5-mercapto-tetrazole, furan, thiophene thiomorpholine possibly optionally mono or di-oxygenated on the S-atom, and amino-cyclohexane possibly substituted by an

hydroxy group/;--

At page 5, lines 11-14, please replace with the the following:

--An even more preferred group of compounds according to the invention are those wherein R, R₁, R₂, R₃, f, m are as above defined and:

R44is a group NR88R99 wherein:

R\$\frac{1}{8} is H or methyl;